AMENDMENTS TO THE CLAIMS

The following listing of claims will replace all prior versions and listings of claims in the application.

LISTING OF CLAIMS

- 1. (Cancelled)
- 2. (Cancelled)
- 3. (Currently Amended) A method for producing an enzyme <u>cellobiase in the presence of glycoslyation inhibitor 2-deoxy-D-glucose from cultures of preparation from a growing culture of Termitomyces clypeatus, said preparation containing high <u>concentration of enzyme increased</u> cellobiase activity in comparison to a control culture, grown in absence of glycoslyation inhibitor 2-deoxy-D-glucose, the <u>said</u> method comprising the steps of:</u>
 - (a) inoculating a mycelial culture of the *Termitomyces clypeatus* into sterilized medium containing <u>carbon and nitrogen sources</u>, <u>inorganic salts</u>, <u>organic nutrients and glycosylation inhibitor 2-deoxy-D-glucose in the range of from about 10 μg/ml to about 2 mg/ml of a glycosylation inhibitor at <u>a</u> pH <u>of</u> between <u>about 3 to 8</u>;</u>
 - (b) growing the mycelial culture at temperatures between 20-37°C under shaking in aerobic conditions; and
 - (c) separating the culture medium from the mycelia to <u>obtain produce</u> the enzyme preparation containing cellobiase activity, <u>said enzyme having an increased enzymatic activity in the range of that is increased at least about 1.15-fold units/ml</u> to about 97 <u>units/ml</u> <u>-fold in the presence of glycosylation inhibitor 2-deoxy-D-glucose</u> in comparison to cellobiase activity produced by the same organism under the same conditions in absence of the glycosylation inhibitor <u>2-deoxy-D-glucose</u>.

- 4. (Cancelled)
- 5. (Cancelled)
- 6. (Previously Presented) The method of claim 3 wherein the medium contains assimilable carbon and nitrogen sources, inorganic salts and organic nutrients.
- 7. (Currently Amended) The method <u>as claimed in claim 3</u>, of claim 6 wherein the <u>assimilable</u> carbon sources <u>of step (a)</u> used are carbohydrates, agrowastes, TCA cycle acids, amino acids, or D-glucosamine wherein the carbohydrates are selected from the group consisting of cellobiose, mannose, fructose, xylose, arabinose, starch, dextrin, cellulose, cotton, and xylan; wherein the agrowastes are selected from the group consisting of baggasse powder, rice-straw powder, wheat bran, corn cob powder, and corn powder; wherein the TCA cycle acids are selected from the group consisting of succinate, fumarate, and maleate; and wherein the amino acids are selected from the group consisting of aspartate, glutamate, serine, histidine, and alanine.
- 8. (Currently Amended) The method of claim 3 wherein the glycosylation inhibitors is of steps (a) and (c) are selected from the group consisting of tunicamycin, 1-deoxynojirimycin, 2-deoxy-D-glucose and D-glucono-lactone.
- 9. (Currently Amended) The method <u>as claimed in claim 3</u>, of claim 6 wherein the <u>assimilable</u> nitrogen source <u>in step (a)</u> is selected from the group consisting of ammonium chloride, ammonium nitrate, ammonium dihydrogen orthophosphate, and potassium nitrate.
- 10. (Currently Amended) The method <u>as claimed in of claim 3</u>, wherein the sterilized medium <u>in step (a)</u> further comprises an organic nutrient selected from the group consisting of malt extract, yeast extract, potato extract, peptone, soya-peptone, bactopeptone, and corn steep liquor.

- 11. (Currently Amended) The method <u>as claimed in</u> of claim 3, wherein the sterilized medium further comprises a detergent selected from the group consisting of Tween-20, Tween-80, and Tween-100.
- 12. (Currently Amended) The method <u>as claimed in claim 3, wherein in the presence of 2-deoxy-D-glucose also enhances activity of other enzymes like endoglucanase and cellobiohydrolase.</u> enzyme preparation containing high cellobiase activity also contains high endo-glucanase activity and high cellobiohydrolase activity.
- 13. (Currently Amended) The method <u>as claimed in</u> of claim 8, wherein <u>enhanced enzyme activity of cellobiase is about 2.23 units/ml in presence of about 0.05mg/ml of 2-deoxy-D-glucose.</u> the onzyme preparation containing high cellobiase activity is an enzyme preparation containing cellobiase activity that is at least about 2.2 units/ml, and wherein the sterilized medium contains about 0.05 mg/ml 2-deoxy-D-glucose.
- 14. (Currently Amended) The method <u>as claimed in of claim 13 8</u>, wherein <u>enhanced enzyme activity of cellobiase is about 50.09 units/ml in presence of about 1 mg/ml of 2-deoxy-D-glucose.</u> the enzyme preparation containing high cellobiase activity is an enzyme preparation having cellobiase activity that is at least about 50 units/ml, wherein the sterilized medium contains about 1 mg/ml 2-deoxy-D-glucose.
- 15. (Currently Amended) The method <u>as claimed in -of claim 14 8</u>, wherein <u>enhanced enzyme activity of cellobiase is</u> the enzyme preparation containing high cellobiase activity is an enzyme preparation having cellobiase activity that is at least about 90 units/ml, wherein the sterilized medium contains about <u>in presence of about 300 µg/ml 2-deoxy-D-glucose</u>.
- 16. (Currently Amended) The method <u>as claimed in</u> of claim 14 <u>8</u>, wherein enhanced enzyme activity of cellobiase is the enzyme preparation containing high cellobiase activity is an enzyme preparation having cellobiase activity that is at least

about 140 units/ml in presence of, wherein the sterilized medium contains about 1 mg/ml 2-deoxy-D-glucose and further contains about 500 µg/ml of 2-deoxy-D-glucose.

- 17. (Currently Amended) The method <u>as claimed in</u> <u>ef-claim 8</u>, wherein <u>enhanced enzyme activity of cellobiase is</u> <u>the enzyme preparation containing high</u> <u>cellobiase activity is an enzyme preparation having cellobiase activity that is at least about 6.18 units/ml in presence of ,wherein the sterilized medium contains at least about 2 mg/ml of glucono-lactone.</u>
- 18. (Currently Amended) The method <u>as claimed in of claim 8</u>, wherein <u>enhanced enzyme activity of cellobiase is</u> the enzyme preparation containing high cellobiase activity is an enzyme preparation having cellobiase activity that is at least about 1.4 units/ml <u>in presence of</u>, wherein the sterilized medium contains at least about 80 µM <u>of</u> 1-deoxynojirimycin.